

chain nodes :

30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46

ring nodes :

1 2 3 4 5 6 7 8 9 10 11 12 13 14 16 17 18 21 22 23 24 25 26

chain bonds :

2-41 2-46 3-42 3-45 4-9 8-33 10-32 11-31 12-30 14-43 14-44 17-39 17-40  
21-37 22-38 24-34 25-35 26-36

ring bonds :

1-2 1-7 1-14 2-3 3-4 4-5 4-14 5-6 6-7 8-9 8-13 9-10 10-11 11-12 12-13  
16-17 16-18 17-18 21-22 21-26 22-23 23-24 24-25 25-26

exact/norm bonds :

1-2 1-7 1-14 4-5 4-14 5-6 6-7 12-30

exact bonds :

2-3 2-41 2-46 3-4 3-42 3-45 4-9 8-33 10-32 11-31 14-43 14-44 16-17  
16-18 17-18 17-39 17-40 21-37 22-38 24-34 25-35 26-36

normalized bonds :

8-9 8-13 9-10 10-11 11-12 12-13 21-22 21-26 22-23 23-24 24-25 25-26

isolated ring systems :

containing 1 : 8 : 16 : 21 :

G1:X, [\*1], [\*2]

Match level :

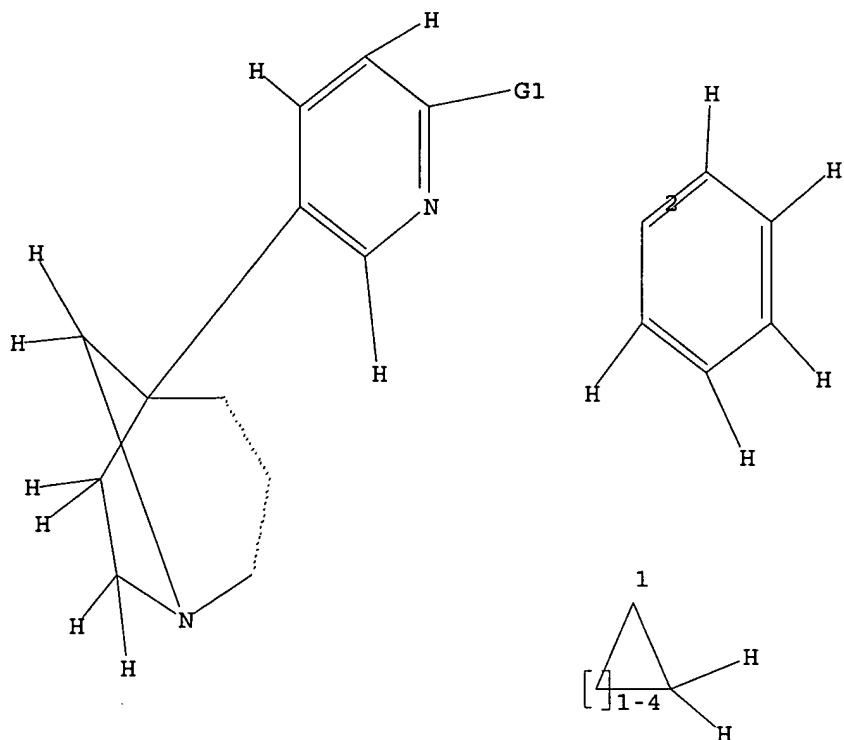
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom  
11:Atom 12:Atom 13:Atom 14:Atom 16:Atom 17:Atom 18:Atom 21:Atom 22:Atom  
23:Atom 24:Atom 25:Atom 26:Atom 30:CLASS 31:CLASS 32:CLASS 33:CLASS  
34:CLASS 35:CLASS 36:CLASS 37:CLASS 38:CLASS 39:CLASS 40:CLASS 41:CLASS  
42:CLASS 43:CLASS 44:CLASS 45:CLASS 46:CLASS

L1 STRUCTURE UPLOADED

=> d

L1 HAS NO ANSWERS

L1 STR



G1 X, [@1], [@2]

Structure attributes must be viewed using STN Express query preparation.

```
=> s 11 ful
FULL SEARCH INITIATED 12:35:59 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 70 TO ITERATE
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100.0% PROCESSED	70 ITERATIONS	10 ANSWERS
SEARCH TIME: 00.00.01		

L2 10 SEA SSS FUL L1

=> fil caplus		SINCE FILE	TOTAL
COST IN U.S. DOLLARS		ENTRY	SESSION
FULL ESTIMATED COST		166.94	167.15

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FILE 'CAPLUS' ENTERED AT 12:36:10 ON 10 FEB 2006
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=> s 12  
L3                1 L2

=> d bib abs

L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2003:534333 CAPLUS

DN 139:101039

TI Derivatives of 5-(pyridin-3-yl)-1-azabicyclo[3.2.1]octane, their preparation, and their application in therapy as nicotinic receptor ligands for treatment of CNS disorders

IN Galli, Frederic; Leclerc, Odile; Lochead, Alistair

PA Sanofi-Synthelabo, Fr.

SO Fr. Demande, 20 pp.

CODEN: FRXBEL

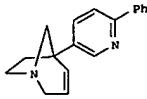
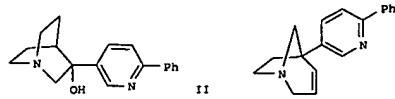
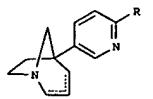
DT Patent

LA French

PAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI FR 2834511	A1	20030711	PR 2002-109	20020107
FR 2834511	B1	20040213		
CA 2471628	AA	20030717	CA 2003-2471628	20030103
WO 2003057697	A1	20030717	WO 2003-FR4	20030103
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW				
AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BP, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2003216777	A1	20030724	AU 2003-216777	20030103
EP 1465893	A1	20041013	EP 2003-712202	20030103
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
BR 2003006707	A	20050209	BR 2003-6707	20030103
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NO 2004002846	A	20040921	NO 2004-2846	20040705
PRAI FR 2002-109	A	20020107		
WO 2003-FR4	W	20030103		
OS MARPAT 139:101039				
GI				

L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2006 ACS on STN (Continued)



AB Title compd. I and their acid addition salts are disclosed [wherein: R = halo, a Ph group (substituted by one or more groups chosen from halo,

C1-6 alkyl or alkoxy, NO<sub>2</sub>, amino, CF<sub>3</sub>, cyano, OH, acetyl, or methylenedioxy], pyridinyl, thiényl, indolyl, or pyrimidinyl (possibly substituted by one or more C1-6 alkoxy); dashed bonds = one single bond and another single

or double bond]. The compds. are useful as pharmaceuticals, particularly as CNS agents, and specifically as ligands of nicotinic receptors. The compds. were tested against nicotinic receptors with the  $\alpha 4\beta 2$  subunit, or with the  $\alpha 7$  subunit. Four synthetic examples and a list of 35 specific compds. (see either di- or tri-HBr or 1:1 oxalate salts)

are given. For instance, 2,5-dibromopyridine was arylated in the 2-position by PhB(OH)<sub>2</sub> using Pd(PPh<sub>3</sub>)<sub>4</sub> catalyst, and the resultant 5-bromo-2-phenylpyridine was lithiated with BuLi and treated with 1-azabicyclo[2.2.1]octan-3-one to give the bicyclic alc. II. Dehydration and rearrangement of II by heating with MeSO<sub>2</sub>H at 180° gave invention compound III, isolated as the di-HBr salt. It tests for

specific binding to isolated rat cerebral nicotinic receptors having either  $\alpha 4\beta 2$  or  $\alpha 7$  subunits, compd. I had IC<sub>50</sub> values in the ranges of 0.01-10  $\mu$ M and 0.005-20  $\mu$ M, resp. Some compds. showed selectivity for the  $\alpha 7$  receptor subtype.

RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT